

1 **Jealous females? Female competition and reproductive suppression in a**  
2 **wild promiscuous primate**

3

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14 **Abstract**

15 Female-female competition over paternal care has rarely been investigated in promiscuous  
16 mammals, where discreet forms of male care have recently been reported despite low  
17 paternity certainty. We investigated female competition over paternal care in a wild  
18 promiscuous primate, the chacma baboon (*Papio ursinus*), where pregnant and lactating  
19 females establish strong social bonds (“friendships”) with males that provide care to their  
20 offspring. We tested whether pregnant and lactating females interfere with the sexual activity  
21 of their male friend to prevent new conceptions that might lead to the subsequent dilution of  
22 his paternal care. We found that pregnant and lactating females were more aggressive towards  
23 oestrous females when they had recently conceived themselves, and when the oestrous female  
24 was mate-guarded by, and showed greater sexual activity with, their male friend. This  
25 aggression also reduced the likelihood of conception of the targeted female. These findings  
26 indicate that females can aggressively prevent further conceptions with their offspring’s carer  
27 through reproductive suppression. Competition over access to paternal care may play an  
28 important and underestimated role in shaping female social relationships and reproductive  
29 strategies in promiscuous mammalian societies.

30

31 **Keywords:** reproductive suppression, paternal care, intrasexual competition, primate, chacma  
32 baboon.

## 33 **1. Introduction**

34 Traditional evolutionary models of paternal care assume that males should only provide care  
35 to their offspring when the degree of paternity certainty is high, typically in monogamous  
36 species [1,2]. Yet recent empirical studies have indicated that male care can also evolve in  
37 promiscuous species where paternity confidence is lower, in particular when the cost of  
38 providing care is also low and does not compromise a male's future reproductive success  
39 [3,4]. For instance, males of several promiscuous primate species provide discreet forms of  
40 care to immatures, such as preferential affiliation, support during conflicts, or tolerance at  
41 feeding sites (e.g. Barbary macaques, *Macaca sylvanus*: [5]; yellow baboons, *Papio*  
42 *cynocephalus*: [6]; olive baboons, *P. anubis*: [7]; black-and-white snub-nosed monkeys,  
43 *Rhinopithecus bieti*: [8]). While males may provide care to unrelated infants to secure future  
44 mating opportunities with the mother [7,9], most studies indicate that males care for their  
45 genetic offspring, and assess their paternity probabilistically based on their mating history  
46 [10–12], or on their offspring's phenotypic resemblance to themselves [13].

47       According to evolutionary theories of parental investment [14], the more offspring a  
48 male sires the more his care will be diluted between them, potentially leading mothers to  
49 compete for exclusive access to their mates [15]. For example, in facultatively polygynous  
50 birds, females breeding with polygynous males experience lower male investment and  
51 reproductive success than those breeding with monogamous males [15–17], and females that  
52 mate first aggressively exclude secondary-mated females from breeding units in order to  
53 maintain male monogamous matings [18]. In polygynous and promiscuous species more  
54 generally, where multiple females mate with a single male that provides paternal services,  
55 females that have already conceived may similarly attempt to prevent further conceptions  
56 with their mate, and the subsequent dilution of paternal care. To do so, they may harass those  
57 females that attempt to mate with him, either to interrupt copulations directly through mating

58 interference and/or to induce chronic physiological stress that reduces their fertility (the  
59 “reproductive suppression hypothesis” [19]).

60         So far, reproductive suppression has been mostly documented in cooperative breeders,  
61 where one or more dominant females use this mechanism to maximise the helper-to-pup ratio  
62 for their own offspring [20–22]. Reproductive suppression has been less well studied in  
63 groups of plural breeders where multiple females reproduce without helpers, but might  
64 similarly represent a manifestation of competition over offspring care provided by males.  
65 Mating interference and female-female harassment are commonplace in plural breeders [23],  
66 and some studies further suggest that the resulting stress can lower the reproductive success of  
67 rivals via physiological mechanisms [24]. For instance, some early studies in yellow baboons  
68 and geladas (*Theropithecus gelada*) indicate that subordinate females that are sexually  
69 receptive are regularly harassed by dominant females, and are also less fertile [25–27].  
70 However, the determinants of female-female competition remain elusive in such studies, as  
71 well as whether the lower fertility of subordinate females is caused by harassment or by other  
72 rank-related differences between females.

73         Here, we tested the hypothesis that females that have already conceived attempt to  
74 prevent new conceptions with the carer of their offspring in a promiscuous primate species,  
75 the chacma baboon (*Papio ursinus*). Chacma baboons live in stable, multimale-multifemale  
76 groups and breed year-round. During pregnancy and lactation, females form a strong social  
77 relationship (‘friendship’) with a particular male [28,29], usually the genetic father of their  
78 offspring [11,12]. Male friends will protect females and their offspring against aggression by  
79 conspecifics [6], which occasionally leads to infanticide [28] and feticide [30], and  
80 subsequently facilitate immature access to ecological resources [31]. Male reproductive skew  
81 is high in chacma baboons [32], which means that high-ranking males are usually involved in  
82 several simultaneous friendships. Female reproductive competition over paternal care appears

83 likely in such societies, as high ranking females aggressively displace subordinates from the  
84 proximity of their male friend [33], and aggression among lactating females peaks in periods  
85 of social instability when infanticide risk is elevated [34]. Moreover, alpha males form weaker  
86 bonds with their offspring than subordinate males [31], suggesting that each offspring indeed  
87 receives less care in the larger paternal sibships of alpha males. We test five predictions of the  
88 reproductive suppression over paternal care hypothesis, namely that pregnant and lactating  
89 females attempt to prevent oestrous females from copulating with their male friend by  
90 harassing them (prediction 1, P1), in particular when they have conceived themselves recently  
91 (P2) (given that synchronous females are expected to compete most intensely over access to  
92 male care); that the intensity of harassment correlates with the fertility (proximity of  
93 ovulation) of the oestrous female (P3) and with the intensity of her sexual activity with the  
94 male friend (P4); and that female-female aggression reduces the probability that the oestrous  
95 female conceives with the male friend (P5).

96

## 97 **2. Material and Methods**

### 98 ***(a) Study site and population***

99 We studied wild chacma baboons at Tsaobis Nature Park, Namibia (22°22'S 15°44'E) (for  
100 details of the site and population, see [35]). We collected data on two habituated groups of  
101 baboons, called 'J' and 'L', over four different periods: June-December 2005, May 2006-  
102 January 2007, June-October 2013 and May-November 2014. Group composition is given in  
103 Table S1 (electronic supplementary materials). Dominance ranks of adult males and females  
104 were established using both *ad libitum* and focal observations of dyadic agonistic interactions  
105 (electronic supplementary materials, Appendix 1).

106

### 107 ***(b) Female reproductive state & mate-guarding patterns***

108 The reproductive state of each female was monitored on a daily basis and categorised as  
109 follows: (1) pregnant, where pregnancy was determined *post hoc* following infant birth, and  
110 encompassed the six months since the conceptive cycle; (2) lactation, if she had a dependant  
111 infant and had not yet resumed cycling, and (3) in oestrus, if she was sexually receptive with a  
112 perineal swelling. Cycling non-swollen females were excluded from the analysis. Every day,  
113 trained observers recorded the swelling state (turgescient or deturgescient) and swelling size of  
114 oestrous females using a semi-quantitative scoring system (from size 1 to 4). For each cycle,  
115 we defined the oestrous period as that time during which a swelling of any size was present,  
116 and the peri-ovulatory period (called hereafter 'POP') as that time during which ovulation  
117 generally occurs, i.e. the 5-day period preceding (and excluding) the day of swelling  
118 detumescence [36]. For each cycle, we determined if it was conceptive or not by identifying *a*  
119 *posteriori* if a pregnancy occurred. The date of conception of pregnant and lactating females  
120 was estimated as the day following detumescence of the conceptive cycle (when witnessed) or  
121 determined *post hoc* by counting back 6 months from the date of birth, the gestation length of  
122 baboons [37]. Mate guarding episodes, defined as periods when oestrous females are  
123 constantly followed by a male that mates exclusively with them and prevents others from  
124 doing so [38], were monitored *ad libitum* on a daily basis.

125

### 126 **(c) Behavioural data**

127 One-hour focal animal samples were conducted on all adult females. In total, our sample  
128 comprises 2971 focal observations on 53 females distributed across reproductive states (Table  
129 S2, electronic supplementary materials). During focal observations, we continuously recorded  
130 aggressive incidents (attacks, chases, threats) and approach-avoid interactions (supplants,  
131 displacements) (for definitions, see Appendix 1, electronic supplementary materials)  
132 involving the focal female, along with the identity of the receiver and initiator. In total, we

133 observed 370 agonistic interactions initiated by pregnant or lactating females toward oestrous  
134 females. In 2013-14, we also recorded all occurrences of male support towards oestrous  
135 females following female aggression. In addition, we recorded every occurrence of male-  
136 female grooming bouts, sexual solicitations ('presenting', when oestrous females present their  
137 hindquarters to males) and copulations, along with the identity of the male. We also noted all  
138 approaches and leaves within 1 meter between the focal individual and any other adult female  
139 (to calculate the time spent in close proximity between female dyads during a focal  
140 observation). Finally, we conducted proximity scans every five minutes to record the identity  
141 and distance of the nearest male neighbour.

142

#### 143 ***(d) Identification of heterosexual friendships***

144 The male friend of each pregnant and lactating female was identified using a combination of  
145 spatial proximity and grooming allocation indices. Full details can be found in Appendix 2 of  
146 the electronic supplementary materials. In short, a male was considered as a friend of a given  
147 pregnant/lactating female if he was both her most frequent nearest neighbour and her most  
148 frequent grooming partner, and if he had an outstandingly high score in both indices  
149 compared to other males (i.e. if his score is at least twice as high as those of other males) [29].  
150 Using this criterion, a female would have either one or two male friend(s) or, in the case of  
151 undifferentiated relationships with males, no friend. Overall, we identified at least one male  
152 friend for 83% of pregnant and lactating females ( $N=67$  out of 81).

153

#### 154 ***(e) Statistical analysis***

155 We ran binomial generalised linear mixed-effect models (GLMMs) using the glmer function  
156 of the lme4 package [39] in R version 3.4.1 [40]. Technical details on how GLMMs were run

157 and how the significance of variables was tested are described in Appendix 3, electronic  
158 supplementary material.

159

160 *Are females more likely to exhibit aggression towards oestrous females that are mate-*  
161 *guarded by their male friend?*

162 We first investigated whether pregnant and lactating females involved in a friendship direct  
163 more aggression towards oestrous females that are mate-guarded by their male friend,  
164 compared to when they are unguarded or mate-guarded by a different male (P1), and when  
165 they are in closer reproductive synchrony, compared to when they are less synchronous (P2).

166 For each focal observation, we created a list of all possible dyadic combinations of initiators  
167 and receivers involving the focal female (e.g., for female A in an entire group including only  
168 two other females, B and C:  $A \rightarrow B$ ,  $A \rightarrow C$ ,  $B \rightarrow A$ ,  $C \rightarrow A$ ). We then restricted this dataset to  
169 dyads where receivers were in oestrous and initiators were pregnant or lactating. We created a  
170 binary variable "Aggression" which recorded whether a directional agonistic interaction  
171 occurred in those dyads during the focal observation. We ran a binomial GLMM using the  
172 occurrence of aggression received by oestrous females from pregnant/lactating females during  
173 a focal observation as the response variable. Random effects comprised the identity of the  
174 initiator and receiver, as well as the identity of the focal observation. The fixed effects  
175 comprised:

- 176 • an index of reproductive synchrony between the initiator and the receiver of the dyad  
177 (in days). This index was calculated as the absolute difference in days between the  
178 initiator's conception date and the focal observation date, and measures the potential  
179 reproductive synchrony should the oestrous female conceive on that focal day.
- 180 • the "mate-guarding status" of the receiver, with three levels: 'Guarded by the friend' if  
181 the receiver was guarded by the male friend of the initiator, 'Guarded by a different



182 male' if the receiver was guarded by another male, and 'Unguarded' if the receiver  
183 was not guarded by any male. Cases where the receiver was guarded by a male and the  
184 initiator had no friend were categorised as 'Guarded by a different male'.

185 • the interaction between the index of potential reproductive synchrony and mate-  
186 guarding status, in order to test whether potential reproductive synchrony is more  
187 important when the receiver is mating with the initiator's friend than in other cases.

188 • an index of spatial proximity between the initiator and the receiver of the dyad (to  
189 control for the fact that two females sharing the same male might attack each other  
190 more often just because they spend more time in proximity). This index was calculated  
191 as the time spent within 1m of each other during the focal observation (calculated  
192 using approaches and leaves within 1m).

193 • the relative dominance rank of the initiator and receiver (as two fixed effects).

194 • group identity (by including group identity as a fixed effect, we do not seek to  
195 systematically assess group effects, which would require a larger sample with targeted  
196 observations, but rather control for such effects should any be present).

197 • year

198

199 *Are females more likely to exhibit aggression towards oestrous females that are closer to*  
200 *conception and copulate more frequently with their male friend?*

201 We tested whether aggression received by an oestrous female from the female friends of a  
202 male (pregnant or lactating) increased with her probability of conception (P3), and the  
203 intensity of her sexual activity with this male (P4), using a binomial GLMM. For each focal  
204 observation of an oestrous female, we identified all possible dyads involving this female and  
205 all resident males that have at least one female friend, and calculated the response variable as  
206 the occurrence of aggression received by that oestrous female from the female friends

207 (pregnant or lactating) of each male (yes/no). Random effects comprised the identity of the  
208 focal female, the male and the focal observation. Fixed factors comprised:

- 209 • an index of fertility measured by proximity from the peri-ovulatory period ('POP'), as  
210 a continuous measure, in days. This measure was used because the probability of  
211 ovulation increases gradually through the oestrus period until reaching a peak in the  
212 five days preceding the day of detumescence, which is easy to identify visually [36].  
213 This index was therefore set to 0 in the 5 days preceding detumescence (i.e. the POP  
214 period), to 1 in the first day preceding the POP, to 2 in the second day preceding the  
215 POP, etc. The day of detumescence was set as 1 (i.e. coded similarly to the first day  
216 preceding the POP period).
- 217 • the rate of sexual activity of the focal female with the male considered (i.e. the number  
218 of presentings and copulations per hour).
- 219 • the mate-guarding status of the oestrous female (guarded by the male/unguarded: 1/0).
- 220 • the interaction between the rate of sexual activity and mate-guarding status, in order to  
221 test whether the effect of sexual activity is more important when the oestrous female is  
222 mate-guarded by the male friend.
- 223 • an index of spatial proximity between the focal female and female friends of a male (to  
224 control for the fact that female friends might attack an oestrous female more often  
225 because she spends more time around them). This index was calculated as the  
226 cumulative time that oestrous females spent within 1m of any female friend of a male  
227 during the focal observation.
- 228 • the relative rank of the oestrous female.
- 229 • the relative rank of the male friend.
- 230 • group identity and year.

231

232

233 *Can females decrease the chance that their male friend conceives with an oestrous female*  
234 *by harassing her?*

235 We then tested whether the probability of conception between an oestrous female and her  
236 mate-guarding male decreased when the oestrous female received higher rates of aggression  
237 from the female friends of the male throughout the oestrus period (P5). Only cycles for which  
238 we had more than four hours of observations of the oestrus period were included (mean±sd  
239 hours of observation per cycle: 16.42±16.42). For each oestrus cycle of each female, we  
240 identified the male that mate-guarded her during her POP (i.e. with whom she may conceive).  
241 For 17 out of 60 cycles, females had several mate-guarding males in her POP; in these cases  
242 we only kept cycles during which one male monopolised 4 days out of 5 of the POP (11/17  
243 cycles) and omitted secondary mate-guarding episodes which were less likely to be  
244 conceptive. For females guarded by males who did not have any female friend, the rate of  
245 aggression was set at zero. We then ran a binomial GLMM using the probability of  
246 conception of each cycle (conceptive/not conceptive: 1/0) as the response variable. Random  
247 effects comprised the identities of the oestrous female and the male.

248 The fixed factors comprised:

- 249 • the rate of aggression received by the oestrous female from the female friends of the  
250 male throughout the oestrus period (calculated as the total number of aggressive  
251 interactions received by the focal female from the female friends of a male throughout  
252 her oestrus cycle, divided by the corresponding observation time).
- 253 • the rate of aggression received by the oestrous female from any other adult female of  
254 the group (calculated as the total number of aggressive interactions received by the  
255 focal female from any non-friend female of a male throughout her oestrus cycle,  
256 divided by the corresponding observation time) to control for a potential confounding

257 effect of female-female aggression at the group level on the chance that the focal  
258 female conceives.

- 259 • whether the cycle was the first postpartum cycle (yes/no) because females experience  
260 reduced fertility in the first cycle following lactational amenorrhea [41])
- 261 • the relative rank of the oestrous female and of the male (over the oestrus period).

262

### 263 **3. Results**

264 Male mating skew was high in both social groups during our study period (see Appendix 4,  
265 electronic supplementary material), and male mating success was highly correlated with male  
266 dominance rank (see Appendix 5). Moreover, resident males had 0 to 9 pregnant and lactating  
267 female friends simultaneously (Table S3), setting-up conditions that may favour female  
268 competition over access to male care.

269

270 *Are females more likely to exhibit aggression towards oestrous females that are mate-*  
271 *guarded by their male friend?*

272 As expected under P1, pregnant and lactating females were more likely to be aggressive  
273 towards oestrous females that were mate-guarded by their male friend (mean dyadic rate±sd:  
274 0.07±0.35 time/h), than towards unguarded females (0.03±0.18) or females guarded by  
275 another male (0.01±0.13) (Table 1, Figure 1a), even when controlling for the fact that females  
276 sharing the same male spend more time in close proximity. Furthermore, pregnant and  
277 lactating females that had conceived more recently were more likely to be aggressive towards  
278 oestrous females (P2) (Table 1, Figure 1b), though this effect of reproductive synchrony was  
279 not greater when the oestrous female was mate-guarded by their male friend than when  
280 unguarded or guarded by another male (interaction between reproductive synchrony and male  
281 sharing status:  $\chi^2_2=1.45$ ,  $p=0.485$ ).

282

283 *Are females more likely to exhibit aggression towards oestrous females that are closer to*  
284 *conception and copulate more frequently with their male friend?*

285 Oestrous females were more likely to receive aggression from the pregnant and lactating  
286 female friends of their sexual partner when they presented to him and copulated with him  
287 more often (P4) (Table 2, Figure 2a). This was true for both mate-guarded and unguarded  
288 females (the interaction between mate-guarding status and the rate of sexual activity was not  
289 significant:  $\chi^2_1=2.63$ ,  $p=0.105$ ), and when controlling for spatial proximity between the  
290 oestrous female and the female friends of her mate. Aggression was also more likely when  
291 the male partner had more female friends, but unaffected by the fertility of the oestrous  
292 female, estimated via her proximity to ovulation (contrary to P3).

293

294 *Can females decrease the chance that their male friend conceives with an oestrous female*  
295 *by harassing her?*

296 Oestrous females received twice as much aggression from the female friends of their mate-  
297 guarding male in non-conceptive cycles (mean±sd amount of aggression received:  $0.13\pm 0.19$ ,  
298  $N=31$  cycles) than in conceptive cycles ( $0.07\pm 0.13$  time/h,  $N=20$  cycles). The probability of  
299 conception of an oestrous female thus decreased when she faced more aggression from the  
300 female friends of her mate (P5), but remained unaffected by aggression received from other  
301 female group-mates (Table 3, Figure 2d). Females were also more likely to conceive with  
302 high-ranking males. The observed association between lower aggression and a greater  
303 likelihood of conception might also arise if mate-guarding males more actively protected  
304 oestrous females during aggressive interactions with other females during conceptive cycles.  
305 However, we assessed the occurrence of male protection of oestrous females (in 2013-14),  
306 and male support was involved in only 9 of 144 aggressive incidences initiated by a pregnant

307 or lactating female towards an oestrous female (including 6 from the mate-guarding male and  
308 3 from other males). Moreover, only one of these cases occurred during a conceptive cycle.  
309 Males therefore rarely intervened in conflicts among females, regardless of their fertility.

310

#### 311 **4. Discussion**

312 High rates of female aggression towards oestrous females have previously been reported in  
313 this [34,42] and other populations of cercopithecids [25–27], raising the question of whether it  
314 represents reproductive suppression. These new analyses extend these studies by showing that  
315 the aggressors include the lactating and pregnant females associated with their male mating  
316 partner (who is the likely father of, and caregiver to, their offspring). This aggression  
317 increases with the sexual activity of the mating couple, and is most likely to occur when the  
318 associated females have conceived more recently and are therefore more vulnerable to the  
319 future dilution of paternal care, especially protection from infanticidal attacks (which are most  
320 common in the first six months of an infant's life [43]). Most importantly, we found that  
321 oestrous females were less likely to conceive during those cycles when they received more  
322 aggression from the female friends of their mate-guarding male. Taken together, these  
323 patterns suggest that females who have already conceived aggressively target oestrous  
324 females who attempt to mate with their offspring's father to prevent him from conceiving  
325 again, which may lead to the loss of paternal services for their own offspring. In mammals,  
326 evidence for reproductive suppression among females primarily comes from cooperative  
327 breeders where dominant females monopolise reproduction to maximise the amount of  
328 allomaternal care received by their offspring. This study reveals a new form of reproductive  
329 suppression in a promiscuous primate society where all females breed and where males  
330 provide discreet paternal care, but the ultimate determinant seems to be essentially similar to  
331 cooperative breeders: to obtain more help to raise offspring.

332 Our interpretation assumes the dilution of paternal care among paternal siblings.  
333 Although this assumption is central to paternal investment theory [14], it may not hold in  
334 promiscuous primates where the cost of paternal care is presumably low, as some forms of  
335 paternal care appear essentially passive. For example, spatial proximity between lactating  
336 females and their male friend is almost exclusively maintained by the female, while the male  
337 simply appears to tolerate their presence [12,28]. However, additional studies suggest that  
338 male care may in fact be more costly than it seems. Playback experiments show that males  
339 readily respond to a distress call from a female friend by running towards her to provide  
340 social support [28] and anecdotal reports show that fathers will engage in severe fights with  
341 rivals that pose an infanticidal threat to their offspring [12]. Fights among adult male baboons  
342 may incur severe to lethal injuries, suggesting that offspring protection can be associated with  
343 life-threatening risks from a male's perspective. It is therefore plausible that males may be  
344 less willing to risk their life when they care for several dependent offspring, if only because  
345 their capacity to protect the remaining offspring will be compromised if they are injured or die  
346 when defending an infant. In line with this, subordinate males, who sire fewer offspring than  
347 dominants, also form closer bonds with their offspring [31]. That said, the paternal care  
348 dilution hypothesis has never been formally tested in promiscuous primates and certainly  
349 deserves further investigation.

350 The exact mechanisms linking female harassment and reproductive suppression in  
351 baboons remain unknown. Reproductive suppression could be mediated by direct mating  
352 interference, given that pregnant and lactating females attack oestrous females more  
353 frequently when they show higher levels of sexual activity. Such aggression may inhibit the  
354 sexual activity of oestrous females, particularly subordinates. However, copulations still occur  
355 at a high frequency during mate-guarding episodes, suggesting that mating interference alone  
356 is unlikely to explain the observed decline in fertility, and that physiological stress may play a

357 critical role. High rates of aggression and elevated levels of cortisol have been found to  
358 disrupt ovulation and the secretion of sex hormones in several captive primates [24,44] and to  
359 cause implantation failure in hamsters [45]. This interpretation is also consistent with our  
360 finding that pregnant and lactating females harass oestrous females not just at the time of  
361 ovulation but throughout the oestrus cycle.

362 Our results suggest that paternal care may be an important determinant of female  
363 competitive relationships in promiscuous primate societies, with wider implications for our  
364 understanding of female reproductive competition across mating systems. In the case of  
365 demography, the ability of some females to suppress synchronous breeding by others may  
366 lead to a staggering of births that could help to explain why some species, like baboons, breed  
367 year-round despite living in seasonal environments [46]. Similarly, the prevalence of sexual  
368 ornaments in oestrous females from promiscuous primate species (e.g. facial colouration [47],  
369 copulatory calls [48] and exaggerated sexual swellings [49]) likely reflects the intensity of  
370 competition faced by females to be chosen by males, despite a typically male-biased sex-ratio.  
371 This study adds new evidence to the idea that these females ultimately compete over access to  
372 male care [50,51].

373

374 **Ethics.** Our research procedures were evaluated and approved by the Ethics  
375 Committee of the Zoological Society of London and the Ministry of Environment and  
376 Tourism (MET), Namibia, and adhered to the ASAB/ABS Guidelines for the Treatment of  
377 Animals in Behavioural Research and Teaching. Our research was conducted under MET  
378 permit numbers 886/2005, 1039/2006, 1786/2013 and 1892/2014.

379

380 **Data accessibility.** Data are available through the public depository GitHub at:  
381 <https://github.com/AliceBaniel/Reproductive-suppression-chacma-baboon>



382

383 **Author's contributions.** A.B. and E.H. designed the study and collected the data. A.B.  
384 performed the analyses. All authors contributed to the drafting of the manuscript.

385

386 **Competing interests.** We have no competing interests.

387

388 **Funding.** A.B. benefitted from financial support from the Agence Nationale de la Recherche  
389 Labex IAST, the Ministère de l'Education Nationale, de l'Enseignement Supérieur et de la  
390 Recherche, the Primate Society of Great Britain, and the Fondation des Treilles.

391

392 **Acknowledgements.** We are grateful to the Tsaobis Baboon Project volunteers in 2005-06  
393 and 2013-14 for invaluable help in the field, and to three anonymous reviewers for very  
394 helpful comments on the manuscript. Thanks to the Tsaobis beneficiaries for permission to  
395 work at Tsaobis Nature Park, the Gobabeb Research and Training Centre for affiliation, the  
396 Ministry of Environment and Tourism for research permits, and to the Snyman and Wittreich  
397 families for permission to work on their land. This paper is a publication of the ZSL Institute  
398 of Zoology's Tsaobis Baboon Project.

399

#### 400 **References**

- 401 1. Kleiman DG, Malcolm JR. 1981 The evolution of male parental investment in  
402 mammals. In *Parental care in mammals* (eds DJ Gubernick, PH Klopfer), pp. 347–387.  
403 New York: Plenum Publishing Corporation.
- 404 2. Clutton-Brock TH. 1991 *The evolution of parental care*. Princeton: Princeton  
405 University Press.
- 406 3. Griffin AS, Alonzo SH, Cornwallis CK. 2013 Why do cuckolded males provide

- 407 paternal care? *PLoS Biol.* **11**.
- 408 4. Alonzo SH, Klug H. 2012 Paternity, maternity, and parental care. In *The evolution of*  
409 *parental care* (eds NJ Royle, PT Smiseth, M Kölliker), pp. 189–205. Oxford: Oxford  
410 University Press.
- 411 5. Paul A, Kuester J, Arnemann J. 1996 The sociobiology of male-infant interactions in  
412 Barbary macaques, *Macaca sylvanus*. *Anim. Behav.* **51**, 155–170.
- 413 6. Buchan JC, Alberts SC, Silk JB, Altmann J. 2003 True paternal care in a multi-male  
414 primate society. *Nature* **425**, 179–181.
- 415 7. Smuts BB. 1985 *Sex and friendship in baboons*. New York: Aldine.
- 416 8. Xiang Z, Sayers K, Grueter C. 2009 Direct paternal care in blackand-white snub-nosed  
417 monkeys. *J. Zool.* **278**, 157–162.
- 418 9. Ménard N, von Segesser F, Scheffrahn W, Pastorini J, Vallet D, Gaci B, Martin RD,  
419 Gautier-Hion A. 2001 Is male-infant caretaking related to paternity and/or mating  
420 activities in wild Barbary macaques (*Macaca sylvanus*). *Comptes Rendus l'Académie*  
421 *des Sci. - Ser. III - Sci. la Vie* **324**, 601–610.
- 422 10. Lehmann J, Fickenscher G, Boesch C. 2007 Kin biased investment in wild  
423 chimpanzees. *Behaviour* **143**, 931–955.
- 424 11. Moscovice LR, Di Fiore A, Crockford C, Kitchen DM, Wittig R, Seyfarth RM, Cheney  
425 DL. 2010 Hedging their bets? Male and female chacma baboons form friendships  
426 based on likelihood of paternity. *Anim. Behav.* **79**, 1007–1015.
- 427 12. Huchard E, Alvergne A, Fejan D, Knapp LA, Cowlshaw G, Raymond M. 2010 More  
428 than friends? Behavioural and genetic aspects of heterosexual associations in wild  
429 chacma baboons. *Behav. Ecol. Sociobiol.* **64**, 769–781.
- 430 13. Widdig A. 2007 Paternal kin discrimination: the evidence and likely mechanisms. *Biol.*  
431 *Rev.* **82**, 319–334.

- 432 14. Trivers RL. 1972 Parental investment and sexual selection. In *Sexual selection and the*  
433 *descent of man* (ed P Campbell), pp. 136–179. London: Heinemann.
- 434 15. Slagsvold T, Lifjeld JT. 1994 Polygyny in birds: the role of competition between  
435 females for male parental care. *Am. Nat.* **143**, 59–94.
- 436 16. Smith HG, Ottosson U, Sandell MI. 1994 Intrasexual competition among polygynously  
437 mated female starlings (*Sturnus vulgaris*). *Behav. Ecol.* **5**, 57–63.
- 438 17. Sandell MI, Smith HG. 1996 Already mated females constrain male mating success in  
439 the European starling. *Proc. R. Soc. B* **263**, 743–747.
- 440 18. Sandell MI. 1998 Female aggression and the maintenance of monogamy: female  
441 behaviour predicts male mating status in European starlings. *Proc. R. Soc. B* **265**,  
442 1307–1311.
- 443 19. Young AJ. 2009 The causes of physiological suppression in vertebrate societies: a  
444 synthesis. In *Reproductive skew in vertebrates: proximate and ultimate causes* (eds R  
445 Hager, CB Jones), pp. 397–436. Cambridge, UK: Cambridge University Press.
- 446 20. Clutton-Brock TH, Hodge SJ, Spong G, Russell AF, Jordan NR, Bennett NC, Sharpe  
447 LL, Manser MB. 2006 Intrasexual competition and sexual selection in cooperative  
448 mammals. *Nature* **444**, 1065–1068.
- 449 21. Clutton-Brock TH, Hodge SJ, P FT, Spong GF, Young AJ. 2010 Adaptive suppression  
450 of subordinate reproduction in cooperative mammals. *Am. Nat.* **176**, 664–673.
- 451 22. Clutton-Brock TH, Russell AF, Sharpe LL, Brotherton PNM, McIlrath GM, White S,  
452 Cameron EZ. 2001 Effects of helpers on juvenile development and survival in  
453 meerkats. *Science* **293**, 2446–2449.
- 454 23. Dixson AF. 1998 *Primate sexuality. Comparative studies of the Prosimians, Monkeys,*  
455 *Apes, and Human beings*. Oxford: Oxford University Press.
- 456 24. Beehner JC, Lu A. 2013 Reproductive suppression in female primates: a review. *Evol.*

- 457 *Anthropol.* **22**, 226–238.
- 458 25. Dunbar RIM. 1980 Determinants and evolutionary consequences of dominance among  
459 female gelada baboons. *Behav. Ecol. Sociobiol.* **7**, 253–265.
- 460 26. Wasser SK, Starling AK. 1988 Proximate and ultimate causes of reproductive  
461 suppression among female yellow baboons at Mikumi National Park, Tanzania. *Am. J.*  
462 *Primatol.* **16**, 97–121.
- 463 27. Wasser SK, Starling AK. 1995 Reproductive competition among female yellow  
464 baboons. In *Primate ontogeny, competition and social behaviour* (eds JG Else, PC  
465 Lee), pp. 343–354. Cambridge: Cambridge University Press.
- 466 28. Palombit RA, Seyfarth RM, Cheney DL. 1997 The adaptive value of ‘friendships’ to  
467 female baboons: experimental and observational evidence. *Anim. Behav.* **54**, 599–614.
- 468 29. Baniel A, Cowlshaw G, Huchard E. 2016 Stability and strength of male-female  
469 associations in a promiscuous primate society. *Behav. Ecol. Sociobiol.* **70**, 761–775.
- 470 30. Zippel MN, Grady JH, Gordon JB, Chow LD, Archie EA, Altmann J, Alberts SC. 2017  
471 Conditional fetal and infant killing by male baboons. *Proc. R. Soc. B* **284**, 20162561.
- 472 31. Huchard E, Charpentier MJ, Marshall H, King AJ, Knapp LA, Cowlshaw G. 2013  
473 Paternal effects on access to resources in a promiscuous primate society. *Behav. Ecol.*  
474 **24**, 229–236.
- 475 32. Weingrill T, Lycett JE, Henzi SP. 2000 Consortship and mating success in chacma  
476 baboons (*Papio cynocephalus ursinus*). *Ethology* **106**, 1033–1044.
- 477 33. Palombit RA, Cheney DL, Seyfarth RM. 2001 Female-female competition for male  
478 ‘friends’ in wild chacma baboons, *Papio cynocephalus ursinus*. *Anim. Behav.* **61**,  
479 1159–1171.
- 480 34. Baniel A, Cowlshaw G, Huchard E. 2018 Context-dependence of female reproductive  
481 competition in wild chacma baboons. *Anim. Behav.* **139**, 37–49.

- 482 35. Cowlshaw G. 1997 Trade-offs between foraging and predation risk determine habitat  
483 use in a desert baboon population. *Anim. Behav.* **53**, 667–686.
- 484 36. Higham JP, Heistermann M, Ross C, Semple S, MacLarnon A. 2008 The timing of  
485 ovulation with respect to sexual swelling detumescence in wild olive baboons.  
486 *Primates* **49**, 295–299.
- 487 37. Altmann J, Altmann S, Hausfater G, McCuskey SA. 1977 Life history of yellow  
488 baboons: physical development, reproductive parameters, and infant mortality.  
489 *Primates* **18**, 315–330.
- 490 38. Alberts SC, Altmann J, Wilson ML. 1996 Mate guarding constrains foraging activity of  
491 male baboons. *Anim. Behav.* **51**, 1269–1277.
- 492 39. Bates D, Maechler M, Bolker B, Walker S. 2014 lme4: linear mixed-effects models  
493 using Eigen and S4. R package version 1.1-7. *R Packag. version 1.1-7*
- 494 40. R Core Team. 2017 R: A language and environment for statistical computing. In *R*  
495 *Foundation for Statistical Computing*, Vienna, Austria.
- 496 41. Gesquiere LR, Wango EO, Alberts S, Altmann J. 2007 Mechanisms of sexual  
497 selection: sexual swellings and estrogen concentrations as fertility indicators and cues  
498 for male consort decisions in wild baboons. *Horm. Behav.* **51**, 114–125.
- 499 42. Huchard E, Cowlshaw G. 2011 Female-female aggression around mating: an extra  
500 cost of sociality in a multimale primate society. *Behav. Ecol.* **22**, 1003–1011.
- 501 43. Palombit RA. 2003 Male infanticide in wild savanna baboons: adaptive significance  
502 and intraspecific variation. In *Sexual selection and reproductive competition in*  
503 *primates: new perspectives and directions* (ed CB Jones), pp. 367–412. American  
504 Society of Primatologists.
- 505 44. Bowman LA, Dilley SR, Keverne EB. 1978 Suppression of oestrogeninduced LH  
506 surges by social subordination in talapoin monkeys. *Nature* **275**, 56–58.

- 507 45. Huck UW, Lisk RD, Miller KS, Bethel A. 1988 Progesterone levels and socially-  
508 induced implantation failure and fetal resorption in golden hamsters (*Mesocricetus*  
509 *auratus*). *Physiol. Behav.* **44**, 321–326.
- 510 46. Ims RA. 1990 The ecology and evolution of reproductive synchrony. *Trends Ecol.*  
511 *Evol.* **5**, 135–140.
- 512 47. Dubuc C, Brent LJM, Accamando AK, Gerald MS, MacLarnon A, Semple S,  
513 Heistermann M, Engelhardt A. 2009 Sexual skin color contains information about the  
514 timing of the fertile phase in free-ranging *Macaca mulatta*. *Int. J. Primatol.* **30**, 777–  
515 789.
- 516 48. O’Connell SM, Cowlishaw G. 1994 Infanticide avoidance, sperm competition and mate  
517 choice: the function of copulation calls in female baboons. *Anim. Behav.* **48**, 687–694.
- 518 49. Nunn CL. 1999 The evolution of exaggerated sexual swellings in primates and the  
519 graded-signal hypothesis. *Anim. Behav.* **58**, 229–246.
- 520 50. Huchard E, Courtiol A, Benavides JA, Knapp LA, Raymond M, Cowlishaw G. 2009  
521 Can fertility signals lead to quality signals? Insights from the evolution of primate  
522 sexual swellings. *Proc. R. Soc. B* **276**, 1889–1897.
- 523 51. Alberts SC, Fitzpatrick CL. 2012 Paternal care and the evolution of exaggerated sexual  
524 swellings in primates. *Behav. Ecol.* **23**, 699–706.

525 **Table 1.** Influence of reproductive synchrony and sharing of the same male partner on the probability of agonistic interactions received by  
 526 oestrous females from pregnant/lactating females. Parameters and tests are based on 2366 focal observations and 276 occurrence of aggressive  
 527 interactions distributed among 50 initiators and 40 receivers GLMMs control for focal observation identity, initiator and receiver identity (fitted  
 528 as random factors). The confidence interval and p-value of statistically significant results are highlighted in bold. SE: Standard error. LRT:  
 529 statistic of a Likelihood Ratio Test. df: degrees of freedom.

530

Response variable	Fixed factor	Levels	Estimate	SE	95% confidence interval	LRT	df	P-value	
Probability that oestrous females receive aggression from the female friends of a male (0/1), by female-female dyad	Reproductive synchrony		-0.45	0.22	<b>[-0.89 ; -0.03]</b>	4.51	1	<b>0.034</b>	
	Male-sharing status	Same male (ref: no male)	0.75	0.18	<b>[0.40 ; 1.10]</b>	38.65	2	<b>&lt;0.001</b>	
		Same male (ref: different male)	1.46	0.24	<b>[0.99 ; 1.94]</b>				
		Different male (ref: no male)	-0.70	0.20	<b>[-1.11 ; -0.32]</b>				
	Spatial proximity		0.24	0.07	<b>[0.08 ; 0.37]</b>	7.91	1	<b>0.005</b>	
	Rank initiator			1.54	0.25	<b>[1.05 ; 2.07]</b>	29.63	1	<b>&lt;0.001</b>
		Rank receiver		-1.06	0.21	<b>[-1.57 ; -0.69]</b>	25.64	1	<b>&lt;0.001</b>
	Group <sup>a</sup>	L		0.57	0.27	<b>[0.03 ; 1.13]</b>	4.27	1	<b>0.039</b>
	Year <sup>b</sup>	2006		0.37	0.35	[-0.30 ; 1.09]	14.38	3	<b>0.002</b>
		2013		-0.70	0.42	[-1.54 ; 0.12]			
2014			0.29	0.36	[-0.43 ; 1.00]				

531

<sup>a</sup> Reference category: J group

532

<sup>b</sup> Reference category: 2005

533

534

535 **Table 2.** Influence of the sexual activity of oestrous females with a male on the probability that they receive aggression from the  
 536 pregnant/lactating females involved in a friendship with him. Parameters and tests are based on 1262 focal observations of oestrous females,  
 537 distributed among 35 focal females and 27 males. We observed 1569 occurrences of sexual activity (587 copulations, 982 presentings), and 199  
 538 occurrences of aggression between oestrous females and pregnant/lactating female friends. GLMMs control for focal observation identity, focal  
 539 female and male identity (fitted as random factors). The confidence interval and p-value of statistically significant results are highlighted in bold.  
 540 SE: Standard error. LRT: statistic of a Likelihood Ratio Test. df: degrees of freedom.

541

Response variable	Fixed factor	Levels	Estimate	SE	95% confidence interval	LRT	df	P-value
Probability that oestrous females receive aggression from the female friends of a male (0/1), across male friends	Sexual activity (presentings, copulations)		0.27	0.13	<b>[0.00 ; 0.53]</b>	3.87	1	<b>0.049</b>
	Mate-guarding with the male <sup>a</sup>		0.71	0.27	<b>[0.18 ; 1.24]</b>	6.83	1	<b>0.009</b>
	Proximity to ovulation		0.23	0.20	[-0.16 ; 0.62]	1.35	1	0.246
	Number of female friends of male		1.32	0.24	<b>[0.86 ; 1.80]</b>	30.76	1	<b>&lt;0.001</b>
	Spatial proximity with female friends		0.21	0.10	[-0.00 ; 0.41]	3.70	1	0.054
	Rank of focal female		-1.03	0.26	<b>[-1.65 ; -0.58]</b>	18.50	1	<b>&lt;0.001</b>
	Rank of male		0.14	0.30	[-0.45 ; 0.74]	0.23	1	0.635
	Group <sup>b</sup>	L	0.73	0.47	[-0.26 ; 1.69]	2.20	1	0.138
	Year <sup>c</sup>	2006	0.34	0.41	[-0.45 ; 1.22]	9.04	3	<b>0.029</b>
	2013	-0.69	0.72	[-2.13 ; 0.75]				
	2014	0.70	0.58	[-0.46 ; 1.91]				

542

<sup>a</sup> Reference category: not mate-guarded by the male

543

<sup>b</sup> Reference category: J group

544

<sup>c</sup> Reference category: 2005

545



546 **Table 3.** Influence of aggression received by oestrous females from the female friends of a male throughout their oestrus periods on the  
 547 probability of conceiving with this male subsequently. Parameters and tests are based on 51 oestrous cycles (out of which 20 were conceptive)  
 548 distributed among 29 focal oestrous females and 18 males. GLMMs control for focal female and male identity (fitted as random factors). The  
 549 confidence interval and p-value of statistically significant results are highlighted in bold. SE: Standard error. LRT: statistic of a Likelihood Ratio  
 550 Test. df: degrees of freedom.

551

Response variable	Fixed factor	Estimate	SE	95% confidence interval	LRT	df	P-value
Probability of conception during a given cycle (0/1)	Aggression received from female friends during oestrus cycle <sup>a</sup>	-1.72	1.02	<b>[-4.06 ; -0.04]</b>	4.07	1	<b>0.044</b>
	Aggression received from other females during oestrus cycle <sup>a</sup>	1.01	0.82	[-0.48 ; 2.84]	1.76	1	0.185
	First postpartum cycle (yes/no)	-0.62	0.82	[-2.36 ; 0.94]	0.59	1	0.441
	Rank of focal female	-0.10	0.74	[-1.59 ; 3.12]	0.02	1	0.889
	Rank of the male	2.65	1.05	<b>[0.91 ; 5.68]</b>	10.51	1	<b>0.001</b>

552 <sup>a</sup> Aggression has been calculated as the number of aggressive interactions that the focal female received throughout her oestrus period  
 553 divided by the corresponding observation time.

554

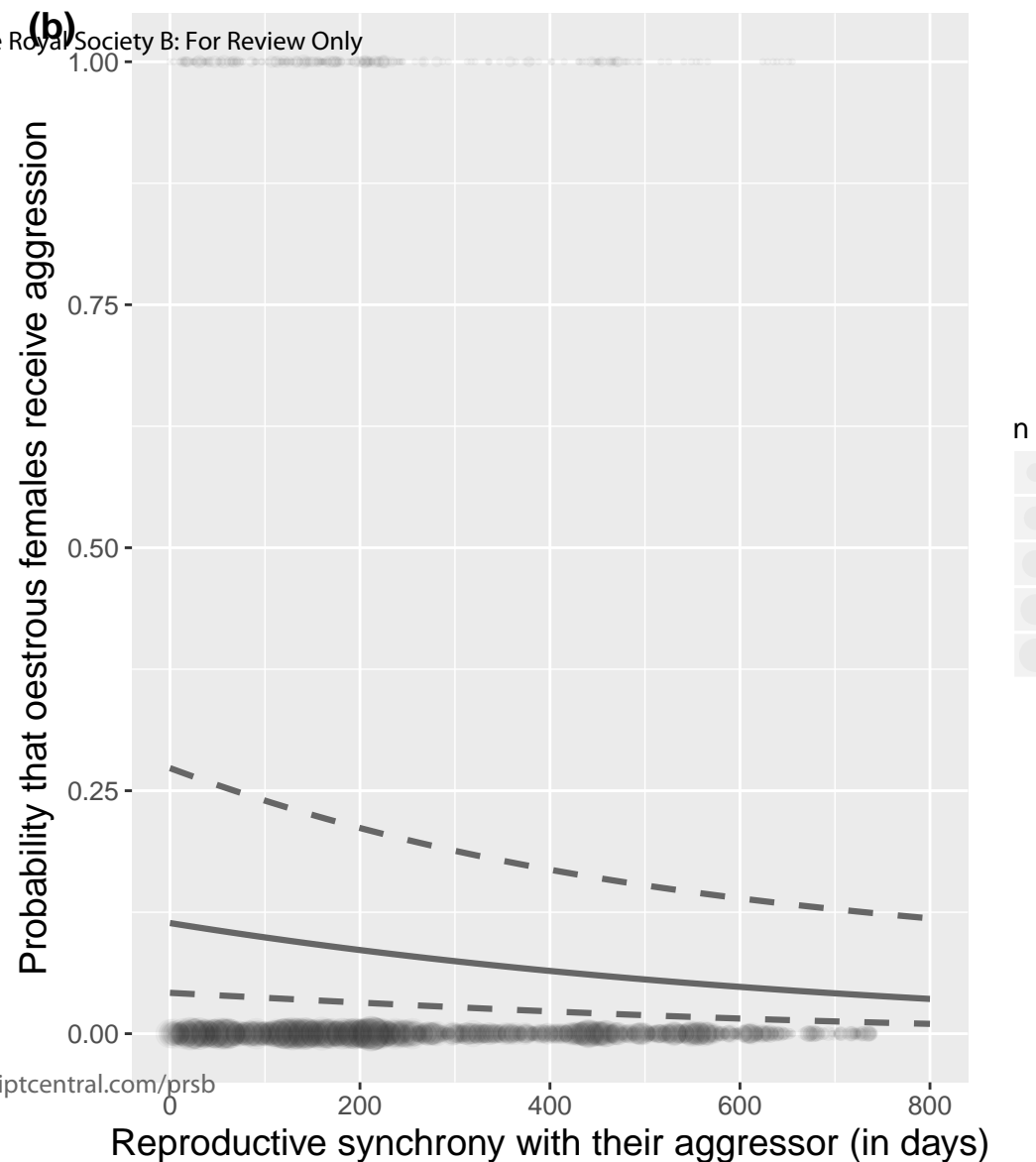
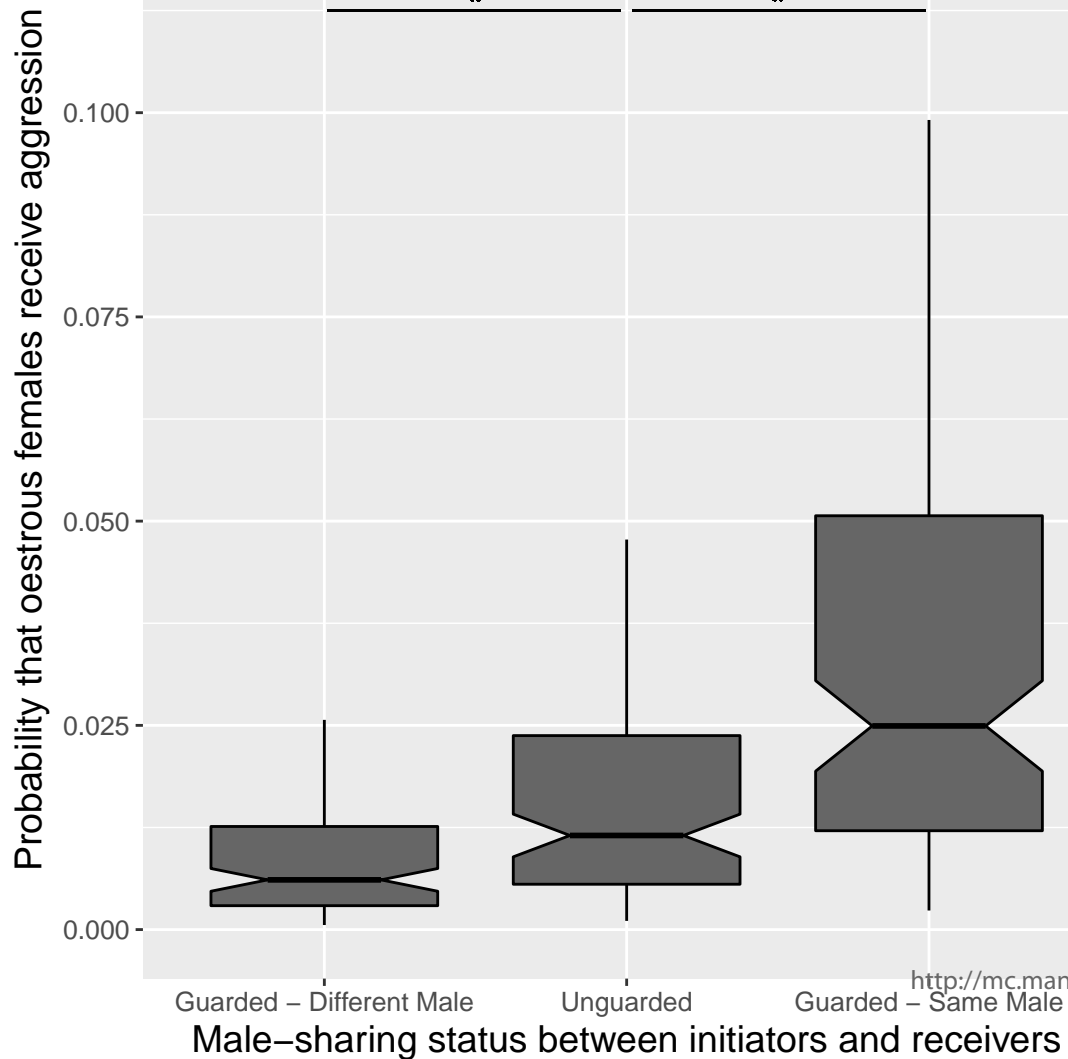
555 **Figures legend**

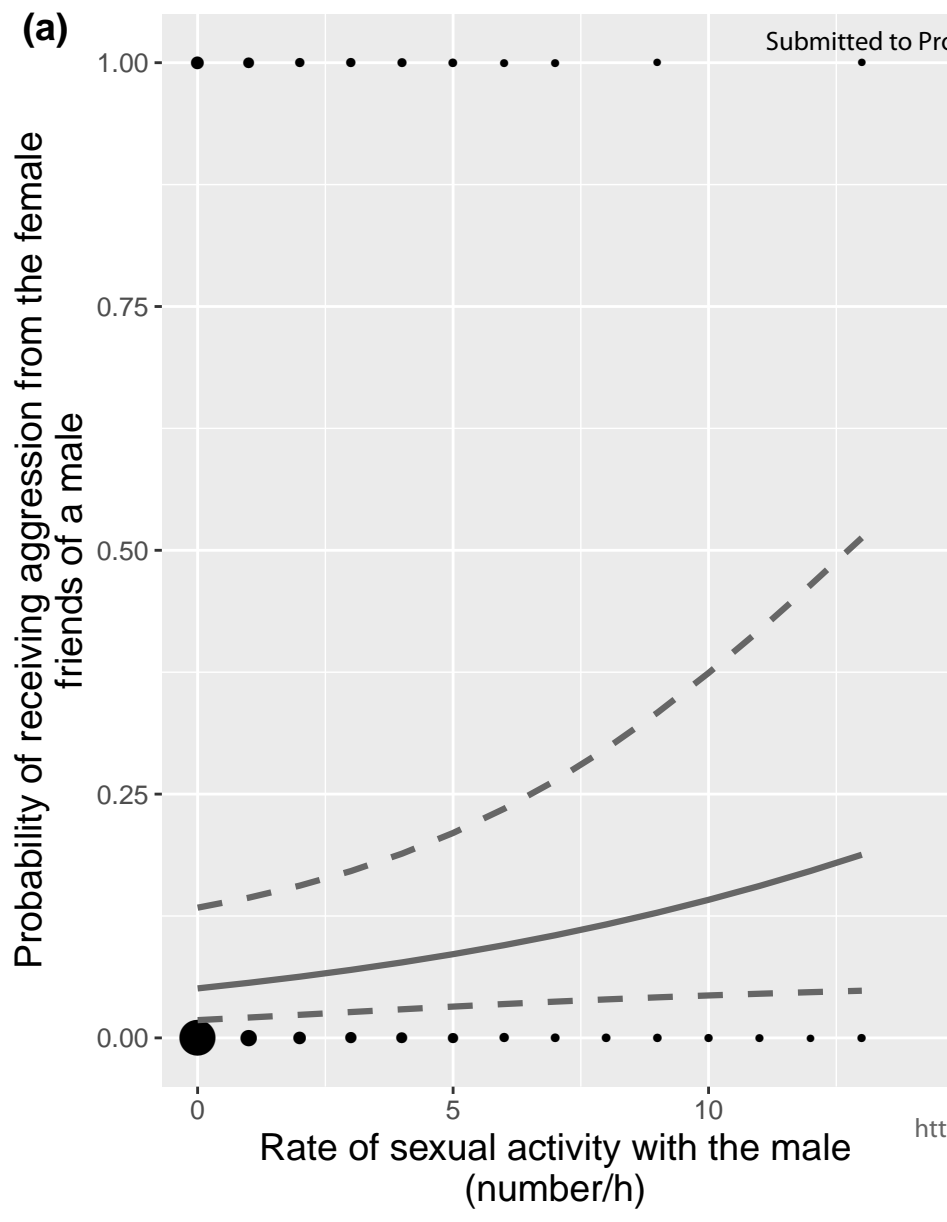
556

557 **Figure 1:** Predicted probability that oestrous females receive aggression from pregnant and  
558 lactating females, according to **(a)** their male sharing status and **(b)** their reproductive  
559 synchrony. In (a), boxplots are drawn from the distribution of the predicted probabilities,  
560 while varying the rank of actors and receivers between 0 and 1, and using the mean for other  
561 numerical values (for a reproductive synchrony of 259 days and a time in proximity of 0.21  
562 minutes), in J troop in 2014 for categorical variables. Comparisons between the different  
563 levels of the variable “male sharing status” are denoted by "\*" if statistically significant. In (b)  
564 the solid line is the model prediction, and the dotted lines represent the 95% confidence  
565 interval. The prediction line is drawn for a dyad sharing a male, that spends 0.21 minutes in  
566 proximity, where the initiator has rank 1 and receiver has rank 0.5, in J group, in 2014. The  
567 grey dots represent the raw data of whether an aggression was exchanged (1) or not (0) within  
568 the dyad, and their size is proportional to the number of occurrences in the dataset.

569

570 **Figure 2:** Predicted probability that oestrous females **(a)** receive aggression from the female  
571 friends of a male, according to their sexual activity with the male and **(b)** conceive with a  
572 male according to the rate of aggression received from the female friends during the oestrus  
573 period. Solid lines represent the model predictions and the dotted lines the 95% confidence  
574 interval. In (a) the prediction line is drawn holding all other fixed effects constant, using the  
575 mean for numeric variables (for a mate-guarded oestrous female of rank 0.53, that spends 0.37  
576 minutes in proximity of the female friends, 7 days before the peri-ovulatory period, and a  
577 male of rank 0.67, having 2.15 female friends), in J group, in 2014 for categorical variables.  
578 In (b) the prediction line is drawn using the mean ranks for female and male (0.57, and 0.79  
579 respectively) and the mean rate of aggression from other females (0.28 aggressive act/hour).  
580 The black dots represent the raw data: whether aggression was exchanged (1) or not (0) in (a),  
581 and whether conception occurred (1) or not (0) in (b), and their size is proportional to the  
582 number of occurrences in the dataset.





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